UK Patent Application (19) GB (11) 2 110 217 A

- (21) Application No 8233945
- (22) Date of filing 29 Nov 1982
- (30) Priority data
- (31) 325174
- (32) 27 Nov 1981
- (33) United States of America
- (43) Application published 15 Jun 1983
- (51) INT CL3
- C08J 9/18 (52) Domestic classification C3C 104 107 110 116 151 153 154 155 180 181 184
 - 358 U1S 1787 C3C
- (56) Documents cited GB 0997356
- (58) Field of search C3C
- (71) Applicants
 USS Engineers and
 Consultants Inc.,
 (USA-Delaware),
 600 Grant Street,
 Pittsburgh,
 State of Pennsylvania
 15230,
- United States of America.
 (72) Inventors
 John James Godfrey;
- Richard Clair Weil.

 (74) Agent and/or Address for Service
 W.P. Thompson and Co., Coopers Building, Church Street, Liverpool,

L1 3AB.

(54) Expandable polystyrene beads

(57) Polyethylene wax of specified physical properties is used as an additive for expandable polystyrene beads to yield small cell size and fast-molding cycles. The polyethylene is substantially linear, has a molecular weight of 700 to 1500 a very narrow molecular-weight distribution (polydispersity of less than 1.2) density at least 15.4 kg/m³ and m.p. of at least 102°C.

GB 2 110 217 A

SPECIFICATION

Expandable polystyrene

5 The present invention relates to expandable polystyrene. Prior to the present invention, it has been known to 5 employ small amounts of polyolefin wax in expandable polystyrene beads in order to achieve more or less uniform cell size and distribution and a relatively rapid pressure drop when the beads are expanded and molded. For example, United States Patent Specifications Nos. 3,224,984 and 3,398,105 disclose the admixture of finely divided polyethylene or polyethylene wax in a concentration of, typically, 0.1 to 0.5 10 percent. The polyethylene disclosed in the aforementioned patents has a molecular weight of 1000 to 4000. It is said that, in the absence of the polyethylene, an undesirable quantity of large cells are formed, causing crystallinity. United States Patent Specification No. 3,060,138 discloses the addition to polystyrene of paraffinic hydrocarbons having 16 to 46 carbon atoms, the paraffinic hydrocarbon being dissolved in an expanding 15 agent, such as, isopentane. Such low-molecular weight materials, however, may be expected to dissolve 15 when contacted by hot liquids, making them undependable for uses such as for coffee cups. United States Patent Specification No. 3,192,169 discloses the use of a paraffin wax which has a melting point of 145°F (63°C) and a density of 0.92 lb/ft³ (14.7kg/m³) at 77°F (25°C), indicating that it contains undesirable non-linearity. It also contains 0.2% oil. It has been found that the use of very specific amounts of polyethylene having very specific characteristics 20 in the manufacture of expandable polystyrene beads provides significant advantages in the subsequent expansion molding of the beads Into articles such as cups. Expanded polystyrene cups are made in large quantities, making the duration of the molding cycle an important economic factor. The uniformity and smallness of the cells are also important. In addition, finished expanded bead cups desirably have to retain 25 their stiffness as well as Impermeability when holding hot liquids such as coffee. 25 In a search for smaller cell size and faster molding cycles, the performance of a series of low-molecularweight homopolymers of ethylene having various molecular weights and narrow molecular-weight distributions was evaluated. It was found that certain polyethylene waxes having linearity (almost no branching) and narrow molecular-weight distribution singularly different from any other commercially 30 available polyethylene were excellent modifiers of polystyrene. The main factors controlling their unique 30 physical properties are: low-molecular weight (that is 700 to 1500), linear molecular structure, and very narrow molecular-weight distribution. The polydispersity (an index of molecular-weight distribution calculated by dividing the weight-average-molecular weight by the number-average molecular weight) is less than 1.2. Due to the absence of side chain branching, plus the narrowness of molecular-weight 35 distribution, these polyethylene waxes differ significantly from other polymers in such physical characteris-35 tics as density, melting point, solubility, melt viscosity, hardness and crystallinity. Their density is at least 0.96 lb/ft³ (15.4kg/m³) and melting point at least 215°F (102°C). Because of the absence of a higher molecular weight fraction, melting point and viscosity are lower than other typical polyethylenes of the same average molecular weight. Also, the absence of very low molecular weight fractions makes the materials harder and permits them to retain their hardness and insolubility at relatively high temperatures. The linearity, or lack of molecular branching, results in a higher density, more crystalline polymer. According, therefore, to the present invention, there is provided a method of making expandable polystyrene beads comprising a polymerizing styrene monomer in a reaction mixture containing 0.05% to 0.5%, by weight based on the styrene, of a polyethylene wax, and impregnating the beads with a blowing 45 agent, said polyethylene wax being substantially linear and having a density of at least 0.96 lb/ft³ 45 (15.4kg/m³), a molecular weight of 700 to 1500, a polydispersity of less than 1.2, and a melting point of at least 215°F (102°C). The invention also provides expandable polystyrene beads containing a blowing agent and 0.05% to about 0.5%, by weight based on the polystyrene, of a polyethylene wax, said polyethylene being substantially 50 linear and having a density of at least 0.96 lb/ft3 (15.4kg/m3), a molecular weight average of 700 to 1500, a 50 polydispersity of less than 1.2, and a melting point of at least 215°F (102°C). Polyethylene wax having a weight average molecular weight of 1000 and a melting point of 235°F (113°C) was most effective (on a weight basis) in reducing cell size; both higher and lower molecular weights were less effective. The existence of an optimum molecular weight for cell size is unknown in the prior art, which 55 generally employs polyethylenes of broader molecular-weight distributions. It was also found that faster 55 molding cycles were obtained as the concentration of additive was increased, up to about 0.5 weight percent. At least about 0.05 weight percent is necessary to produce any appreciable benefit.

Example 1

A suspension polymerization run was conducted in an aqueous suspension with styrene monomer, 0.2
weight percent polyethylene having a weight average molecular weight of 840, a water insoluble
polymerization initiator (peroxide), a suspension agent and a surfactant. The polymer was formed as a slurry
of finely divided beads. These beads were recovered, after acidification to destroy the suspension agent, by
centrifugation. The beads were then dried in a rotary air dryer. The beads were then impregnated with a
blend of n-pentane and isopentane (70-30) at an elevated temperature in an aqueous suspension.

65

2

60

60

Mold Opening

Total cycle time

23 seconds + cooling time

65

	Cooling Component of Cycle				
		Molding Pressure, bars		Cooling Time, seconds	
5	Prior art	2.25		35	5
		2.39		35	
		2.53		55	
10	2.67		55		10
	Invention	2.25	7	7	
15	·	2.39	7	7	15
		2.53	33	· 17	
		2.67	52	37	
20	Example 9 Expandable polystyrene bead	s were made according to	Example 1 but usi	ng 0.2 weight percent 1000	20
	molecular weight polyethylene wax and n-pentane as blowing agent. The beads were pre-expanded in a Rodman-type pre-expander and aged for twenty-four hours in air. An aged density of 1,4 lb/ft ³ (22.4kg/m³)				
26	was obtained. The beads were then transferred to a billet mold (20 in. \times 4 ft. \times 8 ft. (51 cm \times 122 cm \times 244 cm)) and molded at 10 seconds presteam and a maximum of 20 seconds steam (total 30 seconds heading cycle).				1))) 25
23	The following cooling times were obtained:				
	Composition		Cooling Time		
30	Invention		2 min. 56 sec.		30
	Commercial m	aterial	4 min. 50 sec.		
35	Example 10 Expandable polystyrene beads were made according to Example 1. The beads were pre-expanded in a Rodman-type steam pre-expander and aged in air for twenty-four hours. An aged density of 1.3 lb/ft ³ (20.8kg/m ³) was obtained. The beads were then molded at 18 psig (2.25 bars) molding pressure. The over				35
	cycles were as follows:			, motuming pressure. The overa	•••
40	Molding-Cycle	Component	Time, seconds		40
	Mold Closing Filling	•	4 5		
	Steaming Steam Dwell		5 2		
45	Cooling Water Blowout	i.	see below 1		45
	Delay Open Mold Opening		2		
50		Total cycle time 23 s		23 seconds + cooling time	
50	Composition		Cooling Time, sec.	Total Cycle Time, sec.	
	0.1 wt.% 1000	M wt. PE	60	83 .	•
55			60 .	83	55
		5 1000 M wt. PE 5 840 M wt. PE	60	83	
60		5 840 M wt. PE			60
UU	0.2 wt. % 8000	M wt. PE	75	98	
	Prior Art		75	98	

65 M wt. PE = weight average molecular weight polyethylene wax.

15

20

CLAIMS

A method of making expandable polystyrene beads comprising polymerizing styrene monomer in a reaction mixture containing 0.05% to 0.5%, by weight based on the styrene, of a polyethylene wax, and impregnating the beads with a blowing agent, said polyethylene wax being substantially linear and having a density of at least 0.96 lb/ft³ (15.4kg/m³), a molecular weight of 700 to 1500, a polydispersity of less than 1.2, and a melting point of at least 215°F (102°C).

2. A method as claimed in claim 1, in which the polyethylene wax has a molecular weight of 1000 and a melting point of 235°F (113°C).

3. Expandable polystyrene beads containing a blowing agent and 0.05% to about 0.5%, by weight based on the polystyrene, of a polyethylene wax, said polyethylene being substantially linear and having a density of at least 0.96 lb/ft³ (15.4kg/m³), a molecular weight average of 700 to 1500, a polydispersity of less than 1.2, and a melting point of at least 215°F (102°C).

4. Expandable polystyrene beads as claimed in claim 3, in which the polyethylene wax has a molecular weight of 1000 and a melting point of 235°F (113°C).

5. A molded article made from the expandable beads of claim 3 or claim 4.

- 6. A method of making expandable polystyrene beads, as claimed in claim 1, substantially as described in any of Examples 1, 2 and 5 to 10.
- 7. Expandable polystyrene beads as claimd in claim 3 made substantially as described in any of 20 Examples 1, 2 and 5 to 10.

Printed for Her Majesty's Stationery Office, by Croydon Printing Company Limited, Croydon, Surrey, 1983.
Published by The Patent Office, 25 Southampton Buildings, London, WCZA 1AY, from which copies may be obtained.